

WO 96/21421

- 1 -

PCT/FR96/00037

Complete Medium For Use As A Cosmetic And Cosmetic Use Thereof
~~Nutrient medium which can be used as a culture medium~~
~~for epidermal cells and applications~~

The present invention relates to a complex nutrient medium, to its applications and more especially to its use for manufacturing a composition for topical use, and in particular for topical cosmetic or medicinal use.

The composition obtained according to the invention enables an extracellular environment which is entirely suited to the epidermis to be obtained, by supplying in particular:

- an optimized nutritional provision, both in respect of vitamins, ~~and~~ trace elements and ~~in respect of~~ ^{B1 other components} ~~essential amino acids~~
- ~~- cell growth factors directed towards replacing the morphogenic cellular interactions,~~
- and pH and osmolarity characteristics close to physiological conditions.

Generally speaking, according to the invention, the nutritional agent consists of a complex nutrient medium comprising compounds which are both biocompatible, biomimetic and bioavailable in respect of the skin, excluding any biological extract of animal origin, such as foetal calf serum, or of cellular origin.

The complex nutrient medium adopted according to the invention has a composition suitable for permitting, on its own and in an aqueous medium, viable in vitro culture of an inoculum of human epidermal keratinocytes, with at least one clonal proliferation of the latter at the first passage, without a living nourishing substrate such as fibroblasts.

"Biocompatible" is understood to mean the property according to which the compound is harmless to the skin.

"Biomimetic" is understood to mean the fact that the compound is present in the natural state in the skin.

"Bioavailable" is understood to mean the property according to which the compound is assimilable by human epidermal keratinocytes, both in vitro and in vivo.

By routine tests, a person skilled in the art is in a position to formulate a complex nutrient medium according to the invention, in particular by carrying out with the said medium in vitro culturing of keratinocytes, the growth of which can be observed, for example under a microscope.

In this connection, the following documents have already described media suited to in vitro culturing of keratinocytes, the viability and growth of which can be determined by the tests currently in use, and be directly assessed by observation under a microscope:

- Boyce ST, Ham RG, Calcium-regulated differentiation of normal human epidermal keratinocytes in defined clonal culture and serum-free serial culture, J. Invest. Dermatol. 1983; 81: 33S-40S

- Boyce ST, Ham RG, Cultivation, frozen storage, and clonal growth of normal human epidermal keratinocytes ~~in~~ ⁱⁿ ~~serum-free~~ ⁱⁿ serum-free media, J. Tissue Culture Methods. 1985; 9: 83-93.

Where necessary, the content of these publications is incorporated in the present description.

The complex nutrient medium according to the invention comprises amino acids, one or more vitamins, one or more ~~cell growth factors~~ ^{organic components} and one or more inorganic salts.

A composition of the invention for topical use comprises a phase which is biocompatible with the superficial parts of the human body, in which phase at least the said nutrient medium as defined above is distributed homogeneously.

In a composition according to the present invention, the biocompatible phase in which the nutritional agent is distributed can constitute the excipient, or one of the components of the excipient, of the said composition.

Since all of the compounds present in the nutrient medium according to the invention are water-soluble, two methods of formulation may be employed in order to obtain a composition for topical use:

- in the form of an aqueous gel, with the aid of a nonionic water-soluble polymer of the polysaccharide or cellulose ether type (polymers compatible with the high ionic strength of the medium);

10 - in the form of a cosmetic serum.

15 - in emulsified form, on the understanding that
the ionic strength of the discontinuous phase entails
instability of the emulsion; it is, however, possible to
formulate lamellar or cylindrical phases having better
stability, or alternatively a two-phase system re-
emulsified immediately before use by simple shaking;

* in a rigid capsule of the polysaccharide type, dispersed in the lipid phase,

25 The use of liposomes as an encapsulation delivery agent can be envisaged in the form of a liposomal gel in an aqueous continuous phase.

Another subject of the invention is a cosmetic preparation comprising a base defined above, in which the complex nutrient medium constitutes either an active

The complex nutrient medium of the invention can also be used for the preparation or production of a medicament.

A medicinal composition comprising a complex nutrient medium according to the invention can serve as a pharmaceutical formulation base, in particular a nutrient pharmaceutical formulation base.

In addition, a composition as defined above has efficacious properties for preventing or treating disorders of cicatrization such as bedsores, varicose ulcers, stretch marks and keloids, and/or a delay of cicatrization.

7 ? no showing
What
attr no
pamphlets

Example 1 gives an example of a formulation of a composition of the invention.

Example 2 demonstrates the properties of a com-

position of the invention compared to known media, in support of the attached drawing in which:

Fig. 1 is a sectional view of human epidermis after 36 hours of culture in a standard commercial medium designated MCDB 153, marketed, in particular, by IRVINE SCIENTIFIC and GIBCO-BRL,

Fig. 2 is a sectional view of human epidermis after 36 hours of culture in a buffered saline solution (PBS), a balanced saline solution commonly used in cell culture, and

Fig. 3 is a sectional view of human epidermis cultured in the nutrient medium of the invention, described in Example 1, at different culture times:

- A : after 12 hours
- B : after 24 hours
- C : after 36 hours

Example 3 demonstrates the absence of stimulation of the proliferation of transformed cells by a composition of the invention compared to a standard composition, in support of Fig. 4 which depicts a diagram showing the multiplication of transformed cells cultured on a medium of the invention and a standard medium.

Example 4 illustrates the pharmacological properties of a composition of the invention: a) on the treatment of grafts; b) on cicatrization.

Example 1:

Formulation of a composition of the invention

TABLE 1

COMPONENTS	Concentration in mg/l.
30 Amino acids	
L-Alanine	9.2 8.96
L-Arginine HCl <i>HCl</i>	421.4 ✓
L-Asparagine (anhydrous)	14.2 13.2
L-Aspartic acid	4.0 3.99
35 L-Cysteine HCl.H ₂ O	42.0 37.6

2025 RELEASE UNDER E.O. 14176

	Calculated	Found
L-Glutamic acid	14.8	14.71
L-Glutamine	1754.4	877.2
Glycine	7.6	7.51
L-Histidine HCl.H ₂ O	50.0	36.1
5 L-Isoleucine	6.0	33.0
L-Leucine	131.2	132
L-Lysine HCl	54.0	36.6
L-Methionine	13.5	45.0
L-Phenylalanine	10.0	50.0
10 L-Proline	34.6	34.53
L-Serine	126.1	✓
L-Threonine	24.0	23.8
L-Tryptophan	9.3	40.8
L-Tyrosine 2 Na 2H ₂ O	11.7	54.0
15 L-Valine	70.3	70.2
<i>Organic Components</i>		
Vitamins and cell growth factors		
d-Biotin	0.02	0.0146
Folic acid	0.80	0.79
✓ Nicotinamide	0.04	
20 Ca D-Pantothenate	0.30	.285
Pyridoxine HCl	0.06	0.06171
Riboflavin	0.04	0.03764
Thiamine HCl	0.30	0.3373
Vitamin B ₁₂	0.41	0.407
25 i-Inositol	18.0	18.2
Putrescine 2 HCl	0.20	0.1611
Sodium pyruvate	55.0	✓
Thymidine	0.73	0.7266
Adenine (HCl)	24.0	12.16
30 DL-Lipoic acid	0.20	0.2063
Inorganic components		
Sodium chloride	6800.0	6600.0
KCl	112.0	111.83
Na ₂ HPO ₄	284.0	536.2
35 CuSO ₄ .5H ₂ O	0.003	0.00025
Sodium acetate	300.0	(anhydrous) 500.0
D-Glucose	1080.0	1081.0

	HEPES (piperazine)	6600.0	4700.
	Phosphorylethanolamin	0.06768	
	Ethanolamine	0.04684	
	Sodium sulphate	3.4	
5	Sodium bicarbonate	1160.0	1176.0
	FeSO ₄ .7H ₂ O	1.39	1.30
	MgCl ₂ .6H ₂ O	120.0	121.0
	✓ CaCl ₂ .2H ₂ O	from 13.0 to 22.05 14.7	
	ZnSO ₄ .7H ₂ O	0.144	0.1438
10	✓ (NH ₄) ₆ MO ₇ O ₂₄ .4H ₂ O	0.00120	0.00124
	✓ Na ₂ SiO ₃ .5H ₂ O	0.142	0.1421
	✓ MnCl ₂ .4H ₂ O	0.00002	0
	✓ SnCl ₂ .2H ₂ O	0.00011	0.000113
	✓ NH ₄ VO ₃	0.00057	0.00059

15 Example 2:

 The cytocompatibility and the performance features of the complex nutrient medium described in Example 1 were tested on cultures of human keratinocytes in a monolayer, and on human epidermis reconstituted in vitro.

20 The nutrient medium according to Example 1 permits the culture of keratinocytes in a monolayer under optimal conditions of viability for at least 36 hours without the slightest cytotoxic effect manifesting itself.

25 In contrast, a traditional survival solution such as PBS (phosphate buffered saline, a balanced saline solution commonly used in cell culture) proves cytotoxic from 12 hours of incubation onwards.

30 In agreement with Fig. 3, the nutrient medium according to the example permits culture of normal human epidermis reconstituted under optimal conditions of viability, without cytotoxic manifestations even after 36 hours (Fig. 3C) of contact. The cultures displayed basal, prickle, mast and intact, orthokeratotic cornified cell layers, of regular and normal stratification.

35 On comparing Fig. 3C with Fig. 1, the latter

RECEIVED

illustrating the use of a standard commercial medium (MCDB 153, marketed, in particular, by IRVINE SCIENTIFIC), it is seen that the performance features of the medium of the invention are equally good.

5 In contrast, the use of PBS induces, in agreement with Fig. 2, the appearance of keratinocytes in a terminal phase of differentiation at the level of the basal and prickle strata, with more or less pronounced signs of necrosis. A total detachment of the epidermis is
10 also noted, with complete loss of structuring of the different keratinocytic strata.

Example 3: Effects of a composition of the invention on the growth of transformed epidermal cells.

The composition used for this study is the one
15 described in Example 1, comprising the medium termed medium 1.

The effect of the composition 1 on the growth of a spontaneously transformed line of human keratinocytes was tested over 4 days of culture by comparison with
20 cells cultured on a standard medium (DMEM, Dulbecco Modified Epidermal Medium + foetal calf serum).

The cells are first inoculated into the standard medium and grow until the 2nd day after inoculation into this medium. On the 2nd day, the batch of cells is
25 divided into two, one batch continuing to be cultured in standard medium, the other in medium 1.

The results are collated in Figure 4, in which the curve obtained with the points —□— corresponds to the composition of the invention and that obtained with
30 the points --□-- corresponds to the composition of standard medium. The points were duplicated and the counts originate from quadruplicates. The results are corrected for the standard error of the mean, SEM. The arrow seen in the diagram corresponds to the dividing of
35 the batch on the second day of culture.

The morphology of the cells differs according to the medium employed. That of the cells cultured in medium 1 resembles more closely that obtained using a

2025 RELEASE UNDER E.O. 14176

semi-defined medium for epithelial cells, of the GIBCO-BRL KSFM type (cells with looser junctions, less pavement appearance, etc).

5 No significant difference is noted in the growth of this line in accordance with the different media, up to confluence (days 6 to 7, not shown here).

It is concluded that the composition 1 has no stimulatory effect on the proliferation of transformed keratinocytes.

10 Example 4: Effects of a composition of the invention on the taking of human skin grafts and the prevention of cicatrization disorders.

The composition tested is the one described in Example 1, comprising the medium termed medium 1.

15 The effects of the composition 1 on the taking of human skin grafts and the prevention of cicatrization disorders were studied on a mouse model (athymic mouse lacking cell-mediated immunity).

20 Two types of grafts were employed: cultured epidermis and human skins originating from plastic surgery. The grafts were irrigated for 30 days with 1 ml of composition 1 (one application daily) for the group A mice and 1 ml of buffered saline solution (PBS) for the group B mice (20 animals per group). Compresses of tulle gras were applied after each irrigation in order to prevent the grafts from drying out.

25 A clinical observation of the grafts was carried out on D-7, D-15 and D-30.

30 Two parameters were evaluated: the necrosis of the cultured epidermis and the cicatrization.

a) the necrosis of the cultured epidermis ("taking of grafts")

35 Scoring is performed from 0 to 3: 0 = no sign of necrosis; 1 = slight inflammation and superficial degradation of the graft; 2 = partial necrosis; 3 = total necrosis.

The results are collated in Table 2.

TABLE 2

GROUP A MICE

(20 grafts in total, treated with the nutrient composition)

5

Score	D-7	D-15	D-30
0	9/20	12/20	16/20
1	7/20	4/20	0/20
2	3/20	2/20	2/20
3	1/20	2/20	2/20

10

GROUP B MICE

(20 grafts treated with the buffered saline solution)

15

Score	D-7	D-15	D-30
0	2/20	4/20	7/20
1	8/20	6/20	3/20
2	6/20	5/20	5/20
3	4/20	5/20	5/20

20

The composition 1 improves the taking of the grafts of cultured human epidermis on athymic mice compared to a traditional survival solution (PBS). Significant differences are noted from 7 days of treatment onwards, for a final improvement of more than 50%.

b) the cicatrization (with the grafted whole skins)

25

Scoring is performed from 0 to 3: 0 = no cicatrization disorder; 1 = delay of cicatrization; 2 = delay with abnormality of the cicatrization (granulation of the cicatrix); 3 = hypertrophic cicatrix.

The results are collated in Table 3.

TABLE 3

GROUP A MICE

(20 grafted whole skins treated with the nutrient composition)

5

Score	D-7	D-15	D-30
0	20/20	16/20	15/20
1	0/20	3/20	2/20
2	0/20	1/20	2/20
3	0/20	0/20	1/20

10 **GROUP B MICE**

(20 grafted whole skins treated with the buffered saline solution)

15

Score	D-7	D-15	D-30
0	16/20	10/20	5/20
1	4/20	7/20	8/20
2	0/20	3/20	3/20
3	0/20	0/20	4/20

20 The composition 1 significantly improves the cicatrization processes; this effect is especially marked after 30 days of treatment.